



UNITED STATES PATENT AND TRADEMARK OFFICE

UNITED STATES DEPARTMENT OF COMMERCE
United States Patent and Trademark Office
Address: COMMISSIONER FOR PATENTS
P.O. Box 1450
Alexandria, Virginia 22313-1450
www.uspto.gov

APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
10/659,862	09/11/2003	Wael R. Joseph	27839-139 (K-C 19,378C)	5051
45736 7590 04/02/2008 Christopher M. Goff (27839) ARMSTRONG TEASDALE LLP ONE METROPOLITAN SQUARE SUITE 2600 ST. LOUIS, MO 63102				
EXAMINER AHMED, HASAN SYED				
ART UNIT 1618		PAPER NUMBER		
NOTIFICATION DATE 04/02/2008		DELIVERY MODE ELECTRONIC		

Please find below and/or attached an Office communication concerning this application or proceeding.

The time period for reply, if any, is set in the attached communication.

Notice of the Office communication was sent electronically on above-indicated "Notification Date" to the following e-mail address(es):

USpatents@armstrongteasdale.com

Office Action Summary

Application No.

10/659,862

Applicant(s)

JOSEPH ET AL.

Examiner

HASAN S. AHMED

Art Unit

1618

Period for Reply -- The MAILING DATE of this communication appears on the cover sheet with the correspondence address --

A SHORTENED STATUTORY PERIOD FOR REPLY IS SET TO EXPIRE 3 MONTH(S) OR THIRTY (30) DAYS, WHICHEVER IS LONGER, FROM THE MAILING DATE OF THIS COMMUNICATION.

- Extensions of time may be available under the provisions of 37 CFR 1.136(a). In no event, however, may a reply be timely filed after SIX (6) MONTHS from the mailing date of this communication.
- If NO period for reply is specified above, the maximum statutory period will apply and will expire SIX (6) MONTHS from the mailing date of this communication.
- Failure to reply within the set or extended period for reply will, by statute, cause the application to become ABANDONED (35 U.S.C. § 133). Any reply received by the Office later than three months after the mailing date of this communication, even if timely filed, may reduce any earned patent term adjustment. See 37 CFR 1.704(b).

Status

- 1) ☒ Responsive to communication(s) filed on 09 January 2008.
- 2a) ☒ This action is **FINAL**. 2b) ☐ This action is non-final.
- 3) ☐ Since this application is in condition for allowance except for formal matters, prosecution as to the merits is closed in accordance with the practice under *Ex parte Quayle*, 1935 C.D. 11, 453 O.G. 213.

Disposition of Claims

- 4) ☒ Claim(s) 1-14, 16-18 and 20-59 is/are pending in the application.
- 4a) Of the above claim(s) 31-59 is/are withdrawn from consideration.
- 5) ☐ Claim(s) _____ is/are allowed.
- 6) ☒ Claim(s) 1-14, 16-18 and 20-30 is/are rejected.
- 7) ☐ Claim(s) _____ is/are objected to.
- 8) ☐ Claim(s) _____ are subject to restriction and/or election requirement.

Application Papers

- 9) ☐ The specification is objected to by the Examiner.
- 10) ☐ The drawing(s) filed on _____ is/are: a) ☐ accepted or b) ☐ objected to by the Examiner.
Applicant may not request that any objection to the drawing(s) be held in abeyance. See 37 CFR 1.85(a).
Replacement drawing sheet(s) including the correction is required if the drawing(s) is objected to. See 37 CFR 1.121(d).
- 11) ☐ The oath or declaration is objected to by the Examiner. Note the attached Office Action or form PTO-152.

Priority under 35 U.S.C. § 119

- 12) ☐ Acknowledgment is made of a claim for foreign priority under 35 U.S.C. § 119(a)-(d) or (f).
- a) ☐ All b) ☐ Some * c) ☐ None of:
1. ☐ Certified copies of the priority documents have been received.
 2. ☐ Certified copies of the priority documents have been received in Application No. _____.
 3. ☐ Copies of the certified copies of the priority documents have been received in this National Stage application from the International Bureau (PCT Rule 17.2(a)).

* See the attached detailed Office action for a list of the certified copies not received.

Attachment(s)

- 1) ☒ Notice of References Cited (PTO-892)
- 2) ☐ Notice of Draftsperson's Patent Drawing Review (PTO-948)
- 3) ☐ Information Disclosure Statement(s) (PTO/SB/888)
Paper No(s)/Mail Date _____
- 4) ☐ Interview Summary (PTO-413)
Paper No(s)/Mail Date _____
- 5) ☐ Notice of Informal Patent Application
- 6) ☐ Other: _____

DETAILED ACTION

Receipt is acknowledged of applicants' amendment and remarks, which were filed on 9 January 2008.

* * * * *

Claim Rejections - 35 USC § 103

The following is a quotation of 35 U.S.C. 103(a) which forms the basis for all obviousness rejections set forth in this Office action:

(a) A patent may not be obtained though the invention is not identically disclosed or described as set forth in section 102 of this title, if the differences between the subject matter sought to be patented and the prior art are such that the subject matter as a whole would have been obvious at the time the invention was made to a person having ordinary skill in the art to which said subject matter pertains. Patentability shall not be negated by the manner in which the invention was made.

1. Claims 1-14, 16-18, and 20-30 are rejected under 35 U.S.C. 103(a) as being unpatentable over Klofta, *et. al.* (U.S. Patent No. 6,238,682) in view of Krzysik, *et. al.* (U.S. Patent No. 6,440,437 – ('437')) further in view of Krzysik, *et. al.* (U.S. Application No. 2004/0228811 - ('811)).

Klofta, *et. al.* teach a tissue product (see claim 1). The tissue product is comprised of:

- the emollient (fatty acid) of instant claim 1 (see abstract);
- the humectant (polyols) of instant claim 1 (see col. 25, line 16);
- the immobilizing agent (fatty alcohols) of instant claim 1 (see col. 24, lines 4-14);
- the compatibilizing (propylene glycol) agent of instant claim 1 (see col. 17, line 28);
- the fatty acids of instant claim 2 (see abstract);
- the dimethicone of instant claim 3 (see col. 20, line 18);
- the glycerin of instant claims 5-7 (see col. 17, line 21);

Art Unit: 1618

- the polyethylene glycol of instant claims 9 and 10 (*see* col. 17, lines 20-42);
- the stearyl alcohol, of instant claim 11 (*see* col. 24, line 11);
- the propylene glycol of instant claim 12 (*see* col. 17, line 22);
- the dispersing agent of instant claim 13 (*see* col. 22, line 24);
- the polydimethylsiloxanes of instant claim 14 (*see* col. 22, line 24); and
- the surfactant of instant claim 25 (*see* col. 5, line 17).

Klofta, *et. al.* explain that combining the disclosed ingredients into one tissue product is beneficial because they impart, "...a soft and lubricious feel..." *See* col. 4, line 41.

Klofta, *et. al.* teach: (1) about 5% to about 50% emollient (*see* col. 19, lines 25 and 26); (2) about 5% to about 60% humectant (*see* col. 17, line 42); (3) about 5% to about 60% immobilizing agent (*see* col. 27, line 15); and (4) about 5% to about 50% compatibilizing agent (*see* col. 19, lines 25 and 26).

Although Klofta, *et. al.* do not explicitly teach all the percentages recited in instant claims 1, 4, and 8, however, it would have been obvious to one of ordinary skill in the art at the time the invention was made to determine suitable percentages through routine or manipulative experimentation to obtain the best possible results, as these are variable parameters attainable within the art.

Moreover, generally, differences in concentration will not support the patentability of subject matter encompassed by the prior art unless there is evidence indicating such concentration is critical. "[W]here the general conditions of a claim are disclosed in the prior art, it is not inventive to discover the optimum or workable ranges by routine

experimentation.” *In re Aller*, 220 F.2d 454, 456; 105 USPQ 233, 235 (CCPA 1955). Applicants have not demonstrated any unexpected or unusual results, which accrue from the instant percentage ranges.

The Klofta, *et. al.* reference is silent with respect to the (1) phase temperatures of instant claims 1 and 28-30; (2) melting point of instant claim 26; (3) and penetration hardness of instant claim 27. Applicants teach concentration ranges of emollient, humectant, immobilizing agent, and compatibilizing agent that overlap with the prior art. Properties are the same when the structure and composition are the same. Thus, burden shifts to applicant to show unexpected results, by declaration or otherwise. *In re Fitzgerald*, 205 USPQ 594. In the alternative, the claimed properties would have been present once the composition was employed in its intended use. *In re Best*, 195 USPQ 433.

It would have been obvious to a person of ordinary skill in the art at the time the invention was made to combine an emollient, a humectant, an immobilizing agent, and a compatibilizing agent into a tissue product, as taught by Klofta, *et. al.* One of ordinary skill in the art at the time the invention was made would have been motivated to combine these ingredients into a tissue product for the beneficial effect of a soft and lubricious feel, as explained by Klofta, *et. al.*

The Klofta, *et. al.* reference differs from the instant application in that it does not teach the skin barrier of instant claims 15-17, the antioxidant of instant claims 18-20, and the sterol of instant claims 21 and 22.

‘437 teaches a wipe (*see* abstract) comprising:

Art Unit: 1618

- the about 0.1% to about 30% skin barrier enhancing agent of instant claim 1 (see col. 4, line 9);
- the oil of instant claim 16 (see col. 4, line 2);
- the avocado oil of instant claim 17 (see col. 4, line 2);
- 0.3% antioxidant (within the range of instant claim 18; see col. 17, Formulas 1-7);
- the about 0.1% to about 10% sterol of instant claim 21 (see col. 7, line 56); and
- the cholesterol of instant claim 22 (see col. 4, line 4).

'437 explain that combining the disclosed ingredients into one wipe is beneficial because they, "...help maintain skin barrier function..." See col. 2, lines 64-65.

It would have been obvious to a person of ordinary skill in the art at the time the invention was made to combine an emollient, a humectant, an immobilizing agent, a compatibilizing agent, a skin barrier enhancing agent, an antioxidant, and a sterol into a tissue product, as taught by Klofta, *et. al.* in view of '437. One of ordinary skill in the art at the time the invention was made would have been motivated to combine these ingredients into a tissue product for the beneficial effect of a soft and lubricious feel, as explained by Klofta, *et. al.* and to help maintain skin barrier function, as explained by '437

The Klofta, *et. al.* reference differs from the instant application in that it does not teach the butylated hydroxytoluene of instant claims 1 and 20.

'811 teaches a wipe (see abstract) comprising butylated hydroxytoluene (see paragraph 0074).

'811 explains that butylated hydroxytoluene is beneficial because it is an antioxidant and is used to, "...protect (the wipe) from decay or deterioration due to the reaction with oxygen in the air. Antioxidants prevent deterioration which may lead to the generation of rancidity and non-enzymatic browning reaction products." See paragraph 0074.

It would have been obvious to a person of ordinary skill in the art at the time the invention was made to combine an emollient, a humectant, an immobilizing agent, a compatibilizing agent, a skin barrier enhancing agent, an antioxidant such as butylated hydroxytoluene, and a sterol into a tissue product, as taught by Klofta, *et. al.* in view of '437, further in view of '811. One of ordinary skill in the art at the time the invention was made would have been motivated to add butylated hydroxytoluene to a tissue product to prevent deterioration of the tissue, as explained by '811

*

2. Claims 1, 23, and 24 are rejected under 35 U.S.C. 103(a) as being unpatentable over Klofta, *et. al.* (U.S. Patent No. 6,238,682) in view of Krzysik, *et. al.* (U.S. Patent No. 6,440,437 ('437)), further in view of Krzysik, *et. al.* (U.S. Application No. 2004/0228811 - ('811)), further in view of Bowser, *et. al.* (U.S. Patent No. 5,342,976).

Klofta, *et. al.* teach a tissue product (*see above*).

'437 teaches a wipe (*see above*).

'811 teaches a wipe (*see above*).

The Klofta, *et. al.*, '437, and '811 references differ from the instant application in that they do not teach the ceramide and glucosylceramide of instant claims 23 and 24.

Bowser, *et. al.* teach a skin composition that may be used in a tissue product, such as a tissue wipe (*see* col. 16, line 44).

The disclosed composition contains the ceramide and glucosylceramide of instant claims 23 and 24 (*see* col. 1, line 67).

Bowser, *et. al.* explain that a ceramide, such as glucosylceramide, is beneficial in a skin composition because, "...when applied topically to the skin, bring(s) about a marked improvement in skin condition, by enhancing skin barrier function." *See* col. 2, lines 7-9.

It would have been obvious to a person of ordinary skill in the art at the time the invention was made to add a ceramide, such as glucosylceramide to a tissue product, as taught by Klofta, *et. al.* in view of Bowser, *et. al.* One of ordinary skill in the art at the time the invention was made would have been motivated to this ingredient into a tissue product for the beneficial effect of enhancing skin barrier function, as explained by Bowser, *et. al.*

* * * * *

Double Patenting

The nonstatutory double patenting rejection is based on a judicially created doctrine grounded in public policy (a policy reflected in the statute) so as to prevent the unjustified or improper timewise extension of the "right to exclude" granted by a patent and to prevent possible harassment by multiple assignees. A nonstatutory obviousness-type double patenting rejection is appropriate where the conflicting claims are not identical, but at least one examined application claim is not patentably distinct from the reference claim(s) because the examined application claim is either anticipated by, or would have been obvious over, the reference claim(s). *See, e.g., In re Berg*, 140 F.3d 1428, 46 USPQ2d 1226 (Fed. Cir. 1998); *In re Goodman*, 11 F.3d 1046, 29 USPQ2d 1010 (Fed. Cir. 1993); *In re Longi*, 759 F.2d 887, 225 USPQ 645 (Fed. Cir. 1985); *In re Van Ornum*, 686 F.2d 937, 214 USPQ 761 (CCPA 1982); *In re Vogel*, 422

Art Unit: 1618

F.2d 438, 164 USPQ 619 (CCPA 1970); and *In re Thorington*, 418 F.2d 528, 163 USPQ 644 (CCPA 1969).

A timely filed terminal disclaimer in compliance with 37 CFR 1.321(c) or 1.321(d) may be used to overcome an actual or provisional rejection based on a nonstatutory double patenting ground provided the conflicting application or patent either is shown to be commonly owned with this application, or claims an invention made as a result of activities undertaken within the scope of a joint research agreement.

Effective January 1, 1994, a registered attorney or agent of record may sign a terminal disclaimer. A terminal disclaimer signed by the assignee must fully comply with 37 CFR 3.73(b).

Claims 1-14, 16-18, and 20-30 are provisionally rejected on the ground of nonstatutory obviousness-type double patenting as being unpatentable over claims 1-61 of copending Application No. 10/659,969 ('969). Although the conflicting claims are not identical, they are not patentably distinct from each other because '969 claims an absorbent product comprising a moisturizing and lubricating composition comprising an emollient, a humectant, an immobilizing agent, and a compatibilizing agent. See claim 1.

This is a provisional obviousness-type double patenting rejection because the conflicting claims have not in fact been patented.

* * * * *

Response to Arguments

Applicants' arguments filed on 9 January 2008 have been fully considered but they are not persuasive.

1. Applicants argue that neither Klofta nor Krzysik ('437) teach the antioxidants listed in amended claim 1. See remarks, page 24.

Examiner respectfully submits that '437 discloses the antioxidant tocopherol (see col. 17, Formulas 1-7), which was originally listed in instant claims 19 and 20. However,

Art Unit: 1618

applicants have cancelled tocopherol from the claims; as such, the Krzysik ('811) reference has been added to the 35 USC 103 rejection (*see above*) for the proposition that the antioxidant butylated hydroxytoluene was known to be used in tissue products at the time the instant application was filed.

*

2. Applicants argue that Klofta and Krzysik ('437) teach away from each other because Klofta discloses *de minimus* water content (upper limit of 5%) while Krzysik ('437) discloses a higher water content (lower limit of 13.5%) See remarks, pages 27-29. Applicants make the same argument for Klofta and Bowser (lower limit of 15%). See remarks, pages 32-33.

It has been held that a prior art reference must either be in the field of applicant's endeavor or, if not, then be reasonably pertinent to the particular problem with which the applicant was concerned, in order to be relied upon as a basis for rejection of the claimed invention. See *In re Oetiker*, 977 F.2d 1443, 24 USPQ2d 1443 (Fed. Cir. 1992).

In this case, Klofta, Krzysik, and Bowser involve the same field of endeavor, i.e. tissue products. Krzysik was relied upon for the teaching of a skin barrier agent, an antioxidant, and a sterol, while Bowser was relied upon for the teaching of a ceramide. Examiner respectfully submits that all four agents would have the same function in the Klofta tissue product as they do in the Krzysik and Bowser tissue products since the functionality of the three agents does not change between a water content of 5% and a water content of 13.5% or 15%.

* * * * *

Conclusion

Applicants' amendment necessitated the new ground(s) of rejection presented in this Office action. Accordingly, **THIS ACTION IS MADE FINAL**. See MPEP § 706.07(a). Applicant is reminded of the extension of time policy as set forth in 37 CFR 1.136(a).

A shortened statutory period for reply to this final action is set to expire **THREE MONTHS** from the mailing date of this action. In the event a first reply is filed within **TWO MONTHS** of the mailing date of this final action and the advisory action is not mailed until after the end of the **THREE-MONTH** shortened statutory period, then the shortened statutory period will expire on the date the advisory action is mailed, and any extension fee pursuant to 37 CFR 1.136(a) will be calculated from the mailing date of the advisory action. In no event, however, will the statutory period for reply expire later than **SIX MONTHS** from the date of this final action.

★

Correspondence

Any inquiry concerning this communication or earlier communications from the examiner should be directed to **HASAN S. AHMED** whose telephone number is (571)272-4792. The examiner can normally be reached on 9am - 5:30pm.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Michael G. Hartley can be reached on (571)272-0616. The fax phone

Art Unit: 1618

number for the organization where this application or proceeding is assigned is 571-273-8300.

Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see <http://pair-direct.uspto.gov>. Should you have questions on access to the Private PAIR system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free). If you would like assistance from a USPTO Customer Service Representative or access to the automated information system, call 800-786-9199 (IN USA OR CANADA) or 571-272-1000.

/Humera N. Sheikh/
Primary Examiner, Art Unit 1618